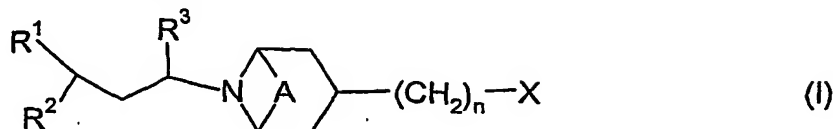


CLAIMS

1. A compound of formula (I):



5 wherein:

A is absent or is  $(CH_2)_2$ ;

$R^1$  is  $C(O)NR^{10}R^{11}$ ,  $C(O)_2R^{12}$ ,  $NR^{13}C(O)R^{14}$ ,  $NR^{15}C(O)NR^{16}R^{17}$ ,  $NR^{18}C(O)_2R^{19}$ , heterocyclyl (for example piperidine, piperazine, pyrrolidine or azetidine), aryl, cycloalkyl or heteroaryl;

10  $R^{10}$ ,  $R^{13}$ ,  $R^{15}$ ,  $R^{16}$  and  $R^{18}$  are hydrogen or  $C_{1-6}$  alkyl;

$R^{11}$ ,  $R^{12}$ ,  $R^{14}$ ,  $R^{17}$  and  $R^{19}$  are  $C_{1-8}$  alkyl (optionally substituted by halo, hydroxy,  $C_{1-6}$  alkoxy,  $C_{1-6}$  haloalkoxy,  $C_{3-6}$  cycloalkyl (optionally substituted by halo),  $C_{5-6}$  cycloalkenyl,  $S(C_{1-4}$  alkyl),  $S(O)(C_{1-4}$  alkyl),  $S(O)_2(C_{1-4}$  alkyl), heteroaryl, aryl, heteroaryloxy or aryloxy), aryl, heteroaryl,  $C_{3-7}$  cycloalkyl (optionally substituted by halo or  $C_{1-4}$  alkyl),  $C_{4-7}$  cycloalkyl fused to a phenyl ring,  $C_{5-7}$  cycloalkenyl, or, heterocyclyl (itself optionally substituted by oxo,  $C(O)(C_{1-6}$  alkyl),  $S(O)_k(C_{1-6}$  alkyl), halo or  $C_{1-4}$  alkyl); or  $R^{11}$ ,  $R^{12}$ ,  $R^{14}$  and  $R^{17}$  can also be hydrogen; or  $R^{10}$  and  $R^{11}$ , and/or  $R^{16}$  and  $R^{17}$  may join to form a 4-, 5- or 6-membered ring which optionally includes a nitrogen, oxygen or sulphur atom, said ring being optionally substituted by  $C_{1-6}$  alkyl,  $S(O)_l(C_{1-6}$  alkyl) or  $C(O)(C_{1-6}$  alkyl);

20  $R^2$  is phenyl, heteroaryl or  $C_{3-7}$  cycloalkyl;

$R^3$  is H or  $C_{1-4}$  alkyl;

X is  $S(O)_2NR^4R^5$  or  $NR^6S(O)_2R^7$ ;

25  $R^7$  is aryl, heteroaryl,  $C_{1-6}$  alkyl,  $C_{3-7}$  cycloalkyl, heterocyclyl or  $NR^8R^9$  wherein  $NR^8R^9$  can be cyclized to form a 4-, 5- or 6-membered ring which optionally includes a nitrogen, oxygen or sulphur atom, said ring being optionally substituted by  $C_{1-6}$  alkyl,  $S(O)_p(C_{1-6}$  alkyl) or  $C(O)(C_{1-6}$  alkyl);

$R^4$  and  $R^8$  are aryl, heteroaryl,  $C_{1-6}$  alkyl (optionally substituted by hydroxy or  $C_{1-6}$  alkoxy),  $C_{3-7}$  cycloalkyl or heterocyclyl;

30  $R^5$ ,  $R^6$  and  $R^9$  are, independently, hydrogen or  $C_{1-6}$  alkyl;

n is 1, 2 or 3;

aryl, phenyl and heteroaryl moieties are independently optionally substituted by one or more of halo, cyano, nitro, hydroxy,  $\text{OC(O)NR}^{20}\text{R}^{21}$ ,  $\text{NR}^{22}\text{R}^{23}$ ,  $\text{NR}^{24}\text{C(O)R}^{25}$ ,  $\text{NR}^{26}\text{C(O)NR}^{27}\text{R}^{28}$ ,  $\text{S(O)}_2\text{NR}^{29}\text{R}^{30}$ ,  $\text{NR}^{31}\text{S(O)}_2\text{R}^{32}$ ,  $\text{C(O)NR}^{33}\text{R}^{34}$ ,  $\text{CO}_2\text{R}^{36}$ ,  $\text{NR}^{37}\text{CO}_2\text{R}^{38}$ ,  $\text{S(O)}_q\text{R}^{39}$ ,  $\text{OS(O)}_2\text{R}^{49}$ ,  $\text{C}_{1-6}$  alkyl (optionally mono-substituted by  $\text{S(O)}_2\text{R}^{50}$  or  $\text{C(O)NR}^{51}\text{R}^{52}$ ),  $\text{C}_{2-6}$  alkenyl,  $\text{C}_{2-6}$  alkynyl,  $\text{C}_{3-10}$  cycloalkyl,  $\text{C}_{1-6}$  haloalkyl,  $\text{C}_{1-6}$  alkoxy( $\text{C}_{1-6}$ )alkyl,  $\text{C}_{1-6}$  alkoxy,  $\text{C}_{1-6}$  haloalkoxy, phenyl, phenyl( $\text{C}_{1-4}$ )alkyl, phenoxy, phenylthio, phenylS(O), phenylS(O)<sub>2</sub>, phenyl( $\text{C}_{1-4}$ )alkoxy, heteroaryl, heteroaryl( $\text{C}_{1-4}$ )alkyl, heteroaryloxy or heteroaryl( $\text{C}_{1-4}$ )alkoxy; wherein any of the immediately foregoing phenyl and heteroaryl moieties are optionally substituted with halo, hydroxy, nitro, S( $\text{C}_{1-4}$  alkyl), S(O)( $\text{C}_{1-4}$  alkyl),  $\text{S(O)}_2(\text{C}_{1-4}$  alkyl),  $\text{S(O)}_2\text{NH}_2$ ,  $\text{S(O)}_2\text{NH}(\text{C}_{1-4}$  alkyl),  $\text{S(O)}_2\text{N}(\text{C}_{1-4}$  alkyl)<sub>2</sub>, cyano,  $\text{C}_{1-4}$  alkyl,  $\text{C}_{1-4}$  alkoxy,  $\text{C(O)NH}_2$ ,  $\text{C(O)NH}(\text{C}_{1-4}$  alkyl),  $\text{C(O)N}(\text{C}_{1-4}$  alkyl)<sub>2</sub>,  $\text{CO}_2\text{H}$ ,  $\text{CO}_2(\text{C}_{1-4}$  alkyl),  $\text{NHC(O)}(\text{C}_{1-4}$  alkyl),  $\text{NHS(O)}_2(\text{C}_{1-4}$  alkyl),  $\text{CF}_3$  or  $\text{OCF}_3$ ;

unless otherwise stated heterocyclyl is optionally substituted by  $\text{C}_{1-6}$  alkyl [optionally substituted by phenyl {which itself optionally substituted by halo,  $\text{C}_{1-4}$  alkyl,  $\text{C}_{1-4}$  alkoxy, cyano, nitro,  $\text{CF}_3$ ,  $\text{OCF}_3$ , ( $\text{C}_{1-4}$  alkyl) $\text{C(O)NH}$ ,  $\text{S(O)}_2\text{NH}_2$ ,  $\text{C}_{1-4}$  alkylthio,  $\text{S(O)}(\text{C}_{1-4}$  alkyl) or  $\text{S(O)}_2(\text{C}_{1-4}$  alkyl)} or heteroaryl {which itself optionally substituted by halo,  $\text{C}_{1-4}$  alkyl,  $\text{C}_{1-4}$  alkoxy, cyano, nitro,  $\text{CF}_3$ , ( $\text{C}_{1-4}$  alkyl) $\text{C(O)NH}$ ,  $\text{S(O)}_2\text{NH}_2$ ,  $\text{C}_{1-4}$  alkylthio,  $\text{S(O)}(\text{C}_{1-4}$  alkyl) or  $\text{S(O)}_2(\text{C}_{1-4}$  alkyl)}], phenyl {optionally substituted by halo,  $\text{C}_{1-4}$  alkyl,  $\text{C}_{1-4}$  alkoxy, cyano, nitro,  $\text{CF}_3$ ,  $\text{OCF}_3$ , ( $\text{C}_{1-4}$  alkyl) $\text{C(O)NH}$ ,  $\text{S(O)}_2\text{NH}_2$ ,  $\text{C}_{1-4}$  alkylthio,  $\text{S(O)}(\text{C}_{1-4}$  alkyl) or  $\text{S(O)}_2(\text{C}_{1-4}$  alkyl)}, heteroaryl {optionally substituted by halo,  $\text{C}_{1-4}$  alkyl,  $\text{C}_{1-4}$  alkoxy, cyano, nitro,  $\text{CF}_3$ , ( $\text{C}_{1-4}$  alkyl) $\text{C(O)NH}$ ,  $\text{S(O)}_2\text{NH}_2$ ,  $\text{C}_{1-4}$  alkylthio,  $\text{S(O)}(\text{C}_{1-4}$  alkyl) or  $\text{S(O)}_2(\text{C}_{1-4}$  alkyl)},  $\text{S(O)}_2\text{NR}^{40}\text{R}^{41}$ ,  $\text{C(O)R}^{42}$ ,  $\text{C(O)}_2(\text{C}_{1-6}$  alkyl) (such as tert-butoxycarbonyl),  $\text{C(O)}_2(\text{phenyl}(\text{C}_{1-2}$  alkyl)) (such as benzyloxycarbonyl),  $\text{C(O)NHR}^{43}$ ,  $\text{S(O)}_2\text{R}^{44}$ ,  $\text{NHS(O)}_2\text{NHR}^{45}$ ,  $\text{NHC(O)R}^{46}$ ,  $\text{NHC(O)NHR}^{47}$  or  $\text{NHS(O)}_2\text{R}^{48}$ , provided none of these last four substituents is linked to a ring nitrogen;

k, l, p and q are, independently, 0, 1 or 2;

$\text{R}^{20}$ ,  $\text{R}^{22}$ ,  $\text{R}^{24}$ ,  $\text{R}^{26}$ ,  $\text{R}^{27}$ ,  $\text{R}^{29}$ ,  $\text{R}^{31}$ ,  $\text{R}^{33}$ ,  $\text{R}^{37}$ ,  $\text{R}^{40}$  and  $\text{R}^{51}$  are, independently, hydrogen or  $\text{C}_{1-6}$  alkyl;

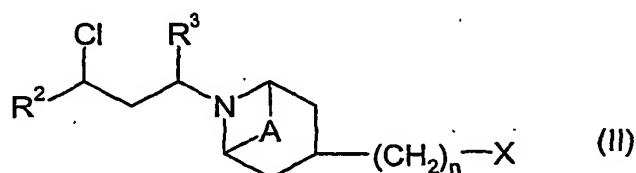
$\text{R}^{21}$ ,  $\text{R}^{23}$ ,  $\text{R}^{25}$ ,  $\text{R}^{28}$ ,  $\text{R}^{30}$ ,  $\text{R}^{32}$ ,  $\text{R}^{34}$ ,  $\text{R}^{36}$ ,  $\text{R}^{38}$ ,  $\text{R}^{39}$ ,  $\text{R}^{41}$ ,  $\text{R}^{42}$ ,  $\text{R}^{43}$ ,  $\text{R}^{44}$ ,  $\text{R}^{45}$ ,  $\text{R}^{46}$ ,  $\text{R}^{47}$ ,  $\text{R}^{48}$ ,  $\text{R}^{49}$ ,  $\text{R}^{50}$  and  $\text{R}^{52}$  are, independently,  $\text{C}_{1-6}$  alkyl (optionally substituted by halo, hydroxy,

C<sub>1-6</sub> alkoxy, C<sub>1-6</sub> haloalkoxy, C<sub>3-6</sub> cycloalkyl, C<sub>5-6</sub> cycloalkenyl, S(C<sub>1-4</sub> alkyl), S(O)(C<sub>1-4</sub> alkyl), S(O)<sub>2</sub>(C<sub>1-4</sub> alkyl), heteroaryl, phenyl, heteroaryloxy or phenyloxy), C<sub>3-7</sub> cycloalkyl, phenyl or heteroaryl; wherein any of the immediately foregoing phenyl and heteroaryl moieties are optionally substituted with halo, hydroxy, nitro, S(C<sub>1-4</sub> alkyl), S(O)(C<sub>1-4</sub> alkyl), S(O)<sub>2</sub>(C<sub>1-4</sub> alkyl), S(O)<sub>2</sub>NH<sub>2</sub>, S(O)<sub>2</sub>NH(C<sub>1-4</sub> alkyl), S(O)<sub>2</sub>N(C<sub>1-4</sub> alkyl)<sub>2</sub>, cyano, C<sub>1-4</sub> alkyl, C<sub>1-4</sub> alkoxy, C(O)NH<sub>2</sub>, C(O)NH(C<sub>1-4</sub> alkyl), C(O)N(C<sub>1-4</sub> alkyl)<sub>2</sub>, CO<sub>2</sub>H, CO<sub>2</sub>(C<sub>1-4</sub> alkyl), NHC(O)(C<sub>1-4</sub> alkyl), NHS(O)<sub>2</sub>(C<sub>1-4</sub> alkyl), C(O)(C<sub>1-4</sub> alkyl), CF<sub>3</sub> or OCF<sub>3</sub>;  
 R<sup>21</sup>, R<sup>23</sup>, R<sup>25</sup>, R<sup>28</sup>, R<sup>30</sup>, R<sup>34</sup>, R<sup>35</sup>, R<sup>36</sup>, R<sup>41</sup>, R<sup>42</sup>, R<sup>43</sup>, R<sup>45</sup>, R<sup>46</sup>, R<sup>47</sup> and R<sup>52</sup> may additionally be hydrogen;  
 or a pharmaceutically acceptable salt thereof or a solvate thereof.

2. A compound as claimed in claim 1 wherein A is absent.
3. A compound as claimed in claim 1 or 2 wherein n is 1 or 2.
4. A compound as claimed in claim 1, 2 or 3 wherein R<sup>3</sup> is hydrogen.
5. A compound as claimed in claim 1, 2, 3 or 4 wherein R<sup>1</sup> is NR<sup>13</sup>C(O)R<sup>14</sup>, wherein R<sup>13</sup> and R<sup>14</sup> are as defined in claim 1.
6. A compound as claimed in claim 1, 2, 3 or 4 wherein R<sup>1</sup> is optionally substituted aryl or optionally substituted heteroaryl, wherein the optional substituents are as recited in claim 1.
7. A compound as claimed in claim 1, 2, 3 or 4 wherein R<sup>1</sup> is optionally substituted heterocyclyl.
8. A compound as claimed in any one of the preceding claims wherein R<sup>2</sup> is phenyl optionally substituted by halo or CF<sub>3</sub>.
9. A compound as claimed in any one of the preceding claims wherein X is NR<sup>6</sup>S(O)<sub>2</sub>R<sup>7</sup>; wherein R<sup>6</sup> and R<sup>7</sup> are as defined in claim 1.

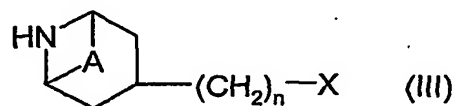
10. A compound as claimed in any one of the preceding claims wherein X is  $S(O)_2NR^4R^5$ ; wherein  $R^4$  and  $R^5$  are as defined in claim 1.

5 11. A process for preparing a compound as claimed in claim 1, the process comprising:  
a. when  $R^1$  is an N-linked optionally substituted heterocycle, reacting a compound of formula (II):

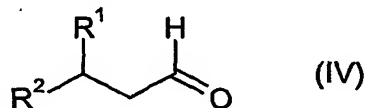


10 wherein  $R^2$ ,  $R^3$ ,  $n$ , A and X are as defined in claim 1, with a compound  $R^1H$  (wherein the H is on a heterocycle ring nitrogen atom) wherein  $R^1$  is as defined above, in the presence of a suitable base, in a suitable solvent and optionally in the presence of sodium iodide;

b. when  $R^3$  is hydrogen, coupling a compound of formula (III):

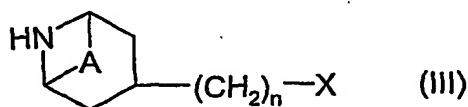


15 wherein  $n$ , A and X are as defined in claim 1, with a compound of formula (IV):



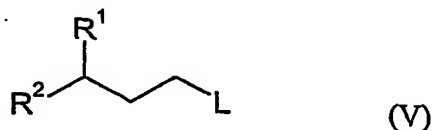
wherein  $R^1$  and  $R^2$  are as defined in claim 1, in the presence of  $NaBH(OAc)_3$  in a suitable solvent at room temperature;

c. when  $R^3$  is hydrogen, coupling a compound of formula (III):



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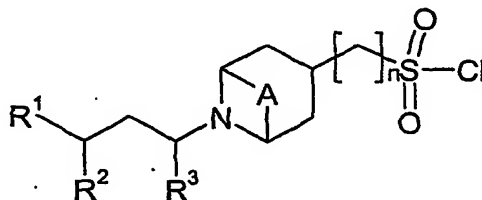
wherein  $n$ , A and X are as defined in claim 1, with a compound of formula (V):



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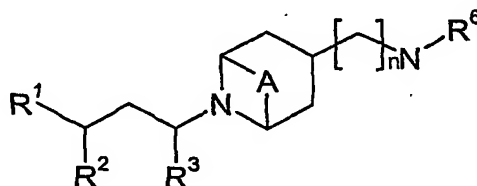
wherein  $R^1$  and  $R^2$  are as defined in claim 1 and L is a leaving group; in the presence of a base, in a suitable solvent at a temperature from 60°C up to the boiling point of the solvent;

d. when X is  $S(O)_2NR^4R^5$ , reacting a compound:



wherein  $R^1$ ,  $R^2$ ,  $R^3$ , A and n are as defined in claim 1, with  $NHR^4R^5$ , wherein  $R^4$  and  $R^5$  are as defined in claim 1, in the presence of a suitable base and in the presence of a suitable solvent; or,

e. when X is  $NR^6S(O)_2NR^7$ , reacting a compound:



wherein  $R^1$ ,  $R^2$ ,  $R^3$ , A and n are as defined in claim 1, with  $R^7S(O)_2Cl$ , in the presence of a suitable base and in the presence of a suitable solvent.

12. A pharmaceutical composition which comprises a compound as claimed in claim 1, or a pharmaceutically acceptable salt thereof or solvate thereof, and a pharmaceutically acceptable adjuvant, diluent or carrier.
13. A compound as claimed in claim 1, or a pharmaceutically acceptable salt thereof or solvate thereof, for use as a medicament.
14. A compound as claimed in claim 1, or a pharmaceutically acceptable salt thereof or solvate thereof, in the manufacture of a medicament for use in therapy.
15. A method of treating a CCR5 mediated disease state comprising administering to a patient in need of such treatment an effective amount of a compound as claimed in claim 1, or a pharmaceutically acceptable salt thereof or solvate thereof.